In the Claims:

Please cancel claims 2, 4, and 5 without prejudice.

Please enter amended claims 1, 3, 6, 8, 10-19, and 21 and add new claim 23 as follows:

1. (once amended) A protease inhibitor represented by the following structure:

 R_{1} -HN R_{2} R_{4} R_{7} R_{6}

wherein

R₁ is selected from the group consisting of hydrogen, carbobenzyloxy-, carbobenzyloxy-glycine-valine-, carbobenzyloxy-alanine-valine-, carbobenzyloxy-leucine-valine-, carbobenzyloxy-phenylalanine-valine-, carbobenzyloxy-serine-valine-, carbobenzyloxy-alanine-asparagine-, carbobenzyloxy-threonine-valine-and carbobenzyloxy-valine-valine-;

R₂ is selected from the group consisting of -CH₂-Phenyl, and -CH₂-CH(CH₃)₂;

R₃ is selected from the group consisting of hydrogen, oxygen and hydroxyl; R₄ is selected from the group consisting of hydrogen, oxygen and hydroxyl, wherein R₃ and R₄ are not both hydroxyl and wherein R₃ and R₄ are either not oxygen or are a single combined oxygen forming a carbonyl group;

 R_5 is selected from the group consisting of hydrogen, and oxygen; R_6 is selected from the group consisting of hydrogen, and oxygen, wherein R_5 and R_6 are either a single combined oxygen forming a carbonyl group or both seperately hydrogen;

R₇ is a radical represented by the formula:

control

NR₈

wherein R_8 is a radical selected from the group consisting of -(H)₂, and -H(t-Butyl); with a proviso that, if either R_2 R_3 is hydroxyl, the R_1 is neither hydrogen nore carbobenzyloxy-.

03

3. (once amended) A stereochemcially pure protease inhibitor represented by the following structure:

wherein

 R_1 is a radical selected from the group consisting of hydrogen, carbobenzyloxy-, carbobenzyloxy-valine-, carbobenzyloxy-glycine-valine-, carbobenzyloxy-alanine-valine-, carbobenzyloxy-leucine-valine-, carbobenzyloxy-phenylalanine-valine-, carbobenzyloxy-serine-valine-, carbobenzyloxy-threonine-valine-, carbobenzyloxy-alanine-asparagine- and carbobenzyloxy-valine-valine-; and

R₂ is a radical selected from the group consisting of -(H)₂, and -H(t-Butyl).

CY

6. (once amended) A protease inhibitor represented by the following structure:

$$R_1$$
-HN $\stackrel{\circ}{\underset{R_2}{\overset{\circ}{=}}}$ R_7

wherein

R₁ is selected from the group consisting of carbobenzyloxy-glycine-valine-, carbobenzyloxy-alanine-valine-, carbobenzyloxy-leucine-valine-, carbobenzyloxy-phenylalanine-valine-, carbobenzyloxy-serine-valine-, carbobenzyloxy-alanine-asparagine-, carbobenzyloxy-threonine-valine- and carbobenzyloxy-valine-valine-;

 R_2 is selected from the group consisting of -CH₂-Phenyl, and -CH₂-CH(CH₃)₂;

 R_7 is a radical represented by the formula:

wherein R₈ is a radical selected from the group consisting of -(H)₂, and -H(t-Butyl).

05

8. (once amended) A protease inhibitor represented by the following structure:

$$R_1$$
-HN $\stackrel{OH}{\underset{\overline{R}_2}{\overset{\circ}{=}}} R_7$

wherein

R₁ is selected from the group consisting of carbobenzyloxy-glycine-valine-, carbobenzyloxy-alanine-valine-, carbobenzyloxy-leucine-valine-, carbobenzyloxy-phenylalanine-valine-, carbobenzyloxy-serine-valine-, carbobenzyloxy-alanine-asparagine-, carbobenzyloxy-threonine-valine- and carbobenzyloxy-valine-valine-;

R₂ is selected from the group consisting of -CH₂-Phenyl, and -CH₂-CH(CH₃)₂;

 R_7 is a radical represented by the formula:



wherein R₈ is a radical selected from the group consisting of -(H)₂, and -H(t-Butyl).

2 Ce

10. (once amended) A stereochemically pure protease inhibitor represented by the following structure:

wherein R_1 is a radical selected from the group consisting of carbobenzyloxy-glycine-valine-, carbobenzyloxy-alanine-valine-, carbobenzyloxy-leucine-valine-, carbobenzyloxy-phenylalanine-valine-, carbobenzyloxy-serine-valine-, carbobenzyloxy-threonine-valine-, carbobenzyloxy-alanine-asparagine- and carbobenzyloxy-valine-valine-; R_2 is selected from the group consisting of $-CH_2$ -Phenyl, and $-CH_2$ -CH(CH_3)₂.

Person 71

11. (once amended) A stereochemically pure protease inhibitor according to claim 10 represented by the following structure:

12. (once amended) A stereochemically pure protease inhibitor according to claim 10 represented by the following structure:

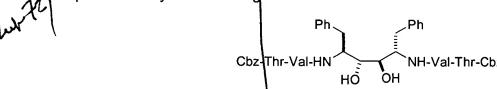
13. (once amended) A stereochemically pure protease inhibitor according to claim 10 represented by the following structure:

Cle

14. (once amended) A stereochemically pure protease inhibitor according to claim 10 represented by the following structure:

15. (once amended) A stereochemically pure protease inhibitor according to claim 10 represented by the following structure:

16. (once amended) A stereochemically pure protease inhibitor according to claim 10 represented by the following structure:



17. (once amended) A stereochemically pure protease inhibitor according to claim 10 represented by the following structure:

18. (once amended)A stereochemically pure protease inhibitor according to claim 3 represented by the following structure:

1 prolate

19. (once amended) A protease inhibitor represented by the following structure:

wherein R_1 is a radical selected from the group consisting of carbobenzyloxy-valine-, carbobenzyloxy-glycine-valine-, carbobenzyloxy-alanine-valine-, carbobenzyloxy-leucine-valine-, carbobenzyloxy-phenylalanine-valine-, carbobenzyloxy-serine-valine-, carbobenzyloxy-threonine-valine-, carbobenzyloxy-valine-valine- and carbobenzyloxy-alanine-asparagine-.

47

21. (once amended) A protease inhibitor represented by the following structure:

wherein R_1 is a radical selected from the group consisting of carbobenzyloxy-valine-, carbobenzyloxy-glycine-valine-, carbobenzyloxy-alanine-valine-, carbobenzyloxy-